

Angelika MÃ¼hlechner

List of Publications by Year in descending order

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Version: 2024-02-01

75
papers

3,177
citations

201674

27
h-index

168389

53
g-index

77
all docs

77
docs citations

77
times ranked

3477
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic pathogenesis of the epileptogenic lesions in Tuberous Sclerosis Complex: Therapeutic targeting of the mTOR pathway. <i>Epilepsy and Behavior</i> , 2022, 131, 107713.	1.7	10
2	Distinct DNA Methylation Patterns of Subependymal Giant Cell Astrocytomas in Tuberous Sclerosis Complex. <i>Cellular and Molecular Neurobiology</i> , 2022, 42, 2863-2892.	3.3	1
3	DNA methylation-based classification of malformations of cortical development in the human brain. <i>Acta Neuropathologica</i> , 2022, 143, 93-104.	7.7	18
4	Down-regulation of the brain-specific cell-adhesion molecule contactin-3 in tuberous sclerosis complex during the early postnatal period. <i>Journal of Neurodevelopmental Disorders</i> , 2022, 14, 8.	3.1	4
5	Increased expression of complement components in tuberous sclerosis complex and focal cortical dysplasia type 2B brain lesions. <i>Epilepsia</i> , 2022, 63, 364-374.	5.1	10
6	Oligosarcomas, IDH-mutant are distinct and aggressive. <i>Acta Neuropathologica</i> , 2022, 143, 263-281.	7.7	18
7	Unexpected Effect of IL-1 β on the Function of GABA _A Receptors in Pediatric Focal Cortical Dysplasia. <i>Brain Sciences</i> , 2022, 12, 807.	2.3	5
8	Upregulation of the pathogenic transcription factor SPI1/PU.1 in tuberous sclerosis complex and focal cortical dysplasia by oxidative stress. <i>Brain Pathology</i> , 2021, 31, e12949.	4.1	11
9	Toward a better definition of focal cortical dysplasia: An iterative histopathological and genetic agreement trial. <i>Epilepsia</i> , 2021, 62, 1416-1428.	5.1	54
10	Balloon cells promote immune system activation in focal cortical dysplasia type 2b. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 826-839.	3.2	14
11	MicroRNA-34a activation in tuberous sclerosis complex during early brain development may lead to impaired corticogenesis. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 796-811.	3.2	5
12	Impaired myelin production due to an intrinsic failure of oligodendrocytes in mTORopathies. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 812-825.	3.2	13
13	Seizure-mediated iron accumulation and dysregulated iron metabolism after status epilepticus and in temporal lobe epilepsy. <i>Acta Neuropathologica</i> , 2021, 142, 729-759.	7.7	31
14	Neurite Outgrowth Inhibitor (NogoA) Is Upregulated in White Matter Lesions of Complex Cortical Malformations. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 274-282.	1.7	0
15	The coding and non-coding transcriptional landscape of subependymal giant cell astrocytomas. <i>Brain</i> , 2020, 143, 131-149.	7.6	24
16	Myelin Pathology Beyond White Matter in Tuberous Sclerosis Complex (TSC) Cortical Tubers. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 1054-1064.	1.7	21
17	Mosaic trisomy of chromosome 1q in human brain tissue associates with unilateral polymicrogyria, very early-onset focal epilepsy, and severe developmental delay. <i>Acta Neuropathologica</i> , 2020, 140, 881-891.	7.7	28
18	Tuberous Sclerosis Complex as Disease Model for Investigating mTOR-Related Gliopathy During Epileptogenesis. <i>Frontiers in Neurology</i> , 2020, 11, 1028.	2.4	25

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19	Reduced expression of the glucocorticoid receptor in the hippocampus of patients with drug-resistant temporal lobe epilepsy and comorbid depression. <i>Epilepsia</i> , 2020, 61, 1595-1605.	5.1	22
20	Seizure outcome and use of antiepileptic drugs after epilepsy surgery according to histopathological diagnosis: a retrospective multicentre cohort study. <i>Lancet Neurology</i> , The, 2020, 19, 748-757.	10.2	177
21	Dysregulation of the MMP/TIMP Proteolytic System in Subependymal Giant Cell Astrocytomas in Patients With Tuberous Sclerosis Complex: Modulation of MMP by MicroRNA-320d In Vitro. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 777-790.	1.7	12
22	Same same but different: A Web-based deep learning application revealed classifying features for the histopathologic distinction of cortical malformations. <i>Epilepsia</i> , 2020, 61, 421-432.	5.1	17
23	Review: Challenges in the histopathological classification of ganglioglioma and DNT: microscopic agreement studies and a preliminary genotype-phenotype analysis. <i>Neuropathology and Applied Neurobiology</i> , 2019, 45, 95-107.	3.2	46
24	Response to ACTH therapy loaded with severe side effects? Do not use synthetic ACTH with the same dosages as "natural" ACTH. <i>Epilepsia</i> , 2019, 60, 1483-1484.	5.1	0
25	In response: The equivalence of the ketogenic diet and adrenocorticotrophic hormone for treatment of infantile spasms: More suggestion than conclusion. <i>Epilepsia</i> , 2019, 60, 2146-2147.	5.1	0
26	Cognitive functioning after epilepsy surgery in children with mild malformation of cortical development and focal cortical dysplasia. <i>Epilepsy and Behavior</i> , 2019, 94, 209-215.	1.7	21
27	Genomic DNA methylation distinguishes subtypes of human focal cortical dysplasia. <i>Epilepsia</i> , 2019, 60, 1091-1103.	5.1	61
28	2017 WONOEP appraisal: Studying epilepsy as a network disease using systems biology approaches. <i>Epilepsia</i> , 2019, 60, 1045-1053.	5.1	12
29	New insights into a spectrum of developmental malformations related to mTOR dysregulations: challenges and perspectives. <i>Journal of Anatomy</i> , 2019, 235, 521-542.	1.5	63
30	Long-term seizure outcome after epilepsy surgery in patients with mild malformation of cortical development and focal cortical dysplasia. <i>Epilepsia Open</i> , 2019, 4, 170-175.	2.4	17
31	Efficacy and tolerability of the ketogenic diet versus high-dose adrenocorticotrophic hormone for infantile spasms: A single-center parallel cohort randomized controlled trial. <i>Epilepsia</i> , 2019, 60, 441-451.	5.1	79
32	Changes in vascular density in resected tissue of 97 patients with mild malformation of cortical development, focal cortical dysplasia or TSC-related cortical tubers. <i>International Journal of Developmental Neuroscience</i> , 2019, 79, 96-104.	1.6	5
33	Single stage epilepsy surgery in children and adolescents with focal cortical dysplasia type II "Prognostic value of the intraoperative electrocorticogram. <i>Clinical Neurophysiology</i> , 2019, 130, 20-24.	1.5	5
34	Oxidative stress and inflammation in a spectrum of epileptogenic cortical malformations: molecular insights into their interdependence. <i>Brain Pathology</i> , 2019, 29, 351-365.	4.1	54
35	Pathophysiology of neurodevelopmental mTOR pathway-associated epileptic conditions: Current status of biomedical research. , 2019, 38, 210-224.		6
36	Ketogenic parenteral nutrition in 17 pediatric patients with epilepsy. <i>Epilepsia Open</i> , 2018, 3, 30-39.	2.4	22

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37	Neuropathology of epilepsy. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 145, 193-216.	1.8	57
38	Epilepsy surgery in infants. Wiener Klinische Wochenschrift, 2018, 130, 341-348.	1.9	10
39	MicroRNA519d and microRNA4758 can identify gangliogliomas from dysembryoplastic neuroepithelial tumours and astrocytomas. Oncotarget, 2018, 9, 28103-28115.	1.8	5
40	Mild Malformation of Cortical Development with Oligodendroglial Hyperplasia in Frontal Lobe Epilepsy: A New Clinicoâ€Pathological Entity. Brain Pathology, 2017, 27, 26-35.	4.1	81
41	Histopathological Findings in Brain Tissue Obtained during Epilepsy Surgery. New England Journal of Medicine, 2017, 377, 1648-1656.	27.0	621
42	Coding and small non-coding transcriptional landscape of tuberous sclerosis complex cortical tubers: implications for pathophysiology and treatment. Scientific Reports, 2017, 7, 8089.	3.3	47
43	Increased expression of (immuno)proteasome subunits during epileptogenesis is attenuated by inhibition of the mammalian target of rapamycin pathway. Epilepsia, 2017, 58, 1462-1472.	5.1	18
44	Impaired oligodendroglial turnover is associated with myelin pathology in focal cortical dysplasia and tuberous sclerosis complex. Brain Pathology, 2017, 27, 770-780.	4.1	51
45	Immediate termination of electrical status epilepticus in sleep after hemispherotomy is associated with significant progress in language development. Developmental Medicine and Child Neurology, 2017, 59, 89-97.	2.1	19
46	Subependymal giant cell astrocytomas in Tuberous Sclerosis Complex have consistent <i>TSC1/TSC2</i> biallelic inactivation, and no <i>BRAF</i> mutations. Oncotarget, 2017, 8, 95516-95529.	1.8	49
47	Characterization of Pathology. , 2017, , 139-160.		5
48	Expression of microRNAs miR21, miR146a, and miR155 in tuberous sclerosis complex cortical tubers and their regulation in human astrocytes and SEGAâ€derived cell cultures. Glia, 2016, 64, 1066-1082.	4.9	51
49	Efficacy and safety of Everolimus in children with TSC - associated epilepsy â€“ Pilot data from an open single-center prospective study. Orphanet Journal of Rare Diseases, 2016, 11, 145.	2.7	47
50	Dysregulation of the (immuno)proteasome pathway in malformations of cortical development. Journal of Neuroinflammation, 2016, 13, 202.	7.2	21
51	Presurgical evaluation of pediatric epilepsy patients prior to hemispherotomy: the prognostic value of 18F-FDG PET. Journal of Neurosurgery: Pediatrics, 2016, 18, 683-688.	1.3	9
52	Functional aspects of early brain development are preserved in tuberous sclerosis complex (TSC) epileptogenic lesions. Neurobiology of Disease, 2016, 95, 93-101.	4.4	50
53	Low-grade epilepsy-associated neuroepithelial tumours â€” the 2016 WHO classification. Nature Reviews Neurology, 2016, 12, 732-740.	10.1	113
54	Specific pattern of maturation and differentiation in the formation of cortical tubers in tuberous sclerosis complex (TSC): evidence from layer-specific marker expression. Journal of Neurodevelopmental Disorders, 2016, 8, 9.	3.1	23

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55	Promoter-Specific Hypomethylation Correlates with IL-1 β Overexpression in Tuberous Sclerosis Complex (TSC). <i>Journal of Molecular Neuroscience</i> , 2016, 59, 464-470.	2.3	23
56	Childhood onset temporal lobe epilepsy: Beyond hippocampal sclerosis. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 228-235.	1.6	7
57	Novel Histopathological Patterns in Cortical Tubers of Epilepsy Surgery Patients with Tuberous Sclerosis Complex. <i>PLoS ONE</i> , 2016, 11, e0157396.	2.5	69
58	Improvement of language development after successful hemispherotomy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2015, 30, 70-75.	2.0	23
59	OP52 " 2694: The ketogenic diet versus ACTH in the treatment of infantile spasms: A prospective randomised study. <i>European Journal of Paediatric Neurology</i> , 2015, 19, S17.	1.6	1
60	OP57 " 2589: Temporal lobe epilepsy in children: Beyond hippocampal sclerosis. <i>European Journal of Paediatric Neurology</i> , 2015, 19, S18-S19.	1.6	0
61	The ketogenic diet in infants " Advantages of early use. <i>Epilepsy Research</i> , 2015, 116, 53-58.	1.6	98
62	Tuberous Sclerosis Complex: new criteria for diagnostic work-up and management. <i>Wiener Klinische Wochenschrift</i> , 2015, 127, 619-630.	1.9	25
63	Efficacy and tolerability of the ketogenic diet in Dravet syndrome " Comparison with various standard antiepileptic drug regimen. <i>Epilepsy Research</i> , 2015, 109, 81-89.	1.6	100
64	7<scp>T MRI</scp> features in control human hippocampus and hippocampal sclerosis: An ex vivo study with histologic correlations. <i>Epilepsia</i> , 2014, 55, 2003-2016.	5.1	76
65	Epilepsy surgery in children and adolescents with malformations of cortical development" Outcome and impact of the new ILAE classification on focal cortical dysplasia. <i>Epilepsy Research</i> , 2014, 108, 1652-1661.	1.6	51
66	Deep sequencing reveals increased DNA methylation in chronic rat epilepsy. <i>Acta Neuropathologica</i> , 2013, 126, 741-756.	7.7	172
67	Vertical perithalamic hemispherotomy: A single-center experience in 40 pediatric patients with epilepsy. <i>Epilepsia</i> , 2013, 54, 1905-1912.	5.1	51
68	Somnolence upon Allergen Provocation in a Child with Hen's Egg Allergy. <i>Klinische Padiatrie</i> , 2013, 225, 232-233.	0.6	1
69	Disconnective surgery in posterior quadrant epilepsy: experience in a consecutive series of 10 patients. <i>Neurosurgical Focus</i> , 2013, 34, E10.	2.3	46
70	Good interobserver and intraobserver agreement in the evaluation of the new ILAE classification of focal cortical dysplasias. <i>Epilepsia</i> , 2012, 53, 1341-1348.	5.1	63
71	Neuropathologic measurements in focal cortical dysplasias: validation of the ILAE 2011 classification system and diagnostic implications for MRI. <i>Acta Neuropathologica</i> , 2012, 123, 259-272.	7.7	106
72	Giant solid-cystic hypothalamic hamartoma. <i>Neurosurgical Focus</i> , 2011, 30, E7.	2.3	14

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73	Neuropathological work-up of focal cortical dysplasias using the new ILAE consensus classification system – practical guideline article invited by the Euro-CNS Research Committee. , 2011, 30, 164-177.		51
74	Use of the ketogenic diet in drug resistant epilepsy syndromes during early infancy: Differences between 3: 1 and 4: 1 formula, a pilot study. Neuropediatrics, 2011, 42, .	0.6	0
75	Beneficial effect of epilepsy surgery in a case of childhood non-paraneoplastic limbic encephalitis. Epilepsy Research, 2010, 90, 295-299.	1.6	12